AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the instant application:

Listing of Claims:

- 1. (Currently amended) A pepsin sensitive modified Cry protein, said protein comprising, characterized in that it has at least one additional pepsin cleavage site relative to a corresponding unmodified Cry protein, wherein said modified Cry protein has increased sensitivity to degradation by pepsin.
- 2. (Currently amended) The modified Cry protein as claimed in claim of Claim 1, characterized in that the wherein the at least one additional pepsin cleavage site is represented by an amino acid residue chosen from leucine, phenylalanine and glutamic acid residues.
- 3. (Currently amended) The modified Cry protein as claimed in either of claims 1 and 2, of Claim 1, characterized in that it is selected from the which is selected from the group consisting of Cry1, Cry3, Cry4, Cry7, Cry8, Cry9, Cry10, Cry16, Cry17, Cry19 and Cry20 proteins.
- 4. (Currently amended) The modified Cry protein as claimed in claim of Claim 3, characterized in that it is a wherein the Cry9 protein is Cry9C protein.
- 5. (Currently amended) The modified Cry protein as claimed in claim of Claim 4, characterized in that it is a wherein the Cry9C protein is Cry9Ca1 protein.
- 6. (Currently amended) The modified Cry protein as claimed in one of claims 1 to 5, characterized in that it has of Claim 1, wherein the at least one additional pepsin cleavage site

is in at least one of the inter-α-helix loops of domain I of said modified Cry protein.

- 7. (Currently amended) The modified Cry protein as claimed in one of claims 1 to 6, characterized in that it has of Claim 6, wherein the at least one additional pepsin cleavage site is in the inter- α -helix loop linking the α 3 and α 4 helices of domain I.
- 8. (Currently amended) The modified Cry protein as claimed in one of claims 5 to 7, characterized in that it has an of Claim 5, wherein the at least one additional pepsin cleavage site is at position 164.
- 9. (Currently amended) The modified Cry protein as claimed in claim 8, characterized in that it is selected from the Cry proteins, the sequences of which are represented by the identifiers of Claim 8, further comprising an amino acid sequence selected from the group consisting of SEQ ID NO:4, SEQ ID NO:6 or and SEQ ID NO:8.
- 10. (Currently amended) The modified Cry protein as claimed in one of claims 1 to 5, characterized in that of Claim 1, wherein the at least one additional pepsin cleavage site is sites are introduced by substituting one or more aspartic acid residues residue with one or more glutamic acid residues residue, substituting one or more tryptophan residues residue with one or more phenylalanine residues residue, and substituting one or more valine or one or more isoleucine residues residue with one or more leucine residues.
- 11. (Currently amended) The modified Cry protein as claimed in claim 11, eharacterized in that of Claim 10, wherein the degree of substitutions which said modified Cry protein possesses is 25% of all aspartic acid, tryptophan, valine or isolecine residues.
- 12. (Currently amended) A method for increasing the pepsin sensitivity of Cry proteins, characterized in that modifying a Cry protein to increase its sensitivity to degradation

by pepsin, comprising introducing into said Cry protein at least one additional pepsin cleavage site is introduced into said Cry proteins relative to a corresponding unmodified Cry protein.

- 13. (Currently amended) The method as claimed in claim 12, characterized in that the of Claim 12, wherein the at least one additional pepsin cleavage site introduced is represented by an amino acid chosen selected from the group consisting of leucine, phenylalanine and glutamic acid-residues.
- 14. (Currently amended) The method as claimed in either of claims 12 and 13, characterized in that it applies to the Cry proteins of Claim 12, wherein the Cry protein to be modified is selected from the group consisting of Cry1, Cry3, Cry4, Cry7, Cry8, Cry9, Cry10, Cry16, Cry17, Cry19 and Cry20 proteins.
- 15. (Currently amended) The method as claimed in claim 14, characterized in that it applies to of Claim 14, wherein the Cry9 protein is the Cry9C protein.
- 16. (Currently amended) The method as claimed in claim 15, characterized in that it applies to of Claim 15, wherein the Cry9C protein is the Cry9Ca1 protein.
- 17. (Currently amended) The method as claimed in one of claims 12 to 16, characterized in that of Claim 12, wherein the at least one additional pepsin cleavage site is introduced into at least one of the inter-α-helix loops of domain I of said modified Cry proteins protein.
- 18. (Currently amended) The method as claimed in one of claims 12 to 17, eharacterized in that of Claim 17, wherein the at least one additional pepsin cleavage site is introduced into the inter-α-helix loop linking the α and α4 helices of domain I of said modified Cry protein.

- 19. (Currently amended) The method as claimed in one of claims 16 to 18, characterized in that an of Claim 16, wherein the at least one additional pepsin cleavage site is introduced at position 164.
- 20. (Currently amended) The method as claimed in one of claims 12 to 16, characterized in that the of Claim 12, where the at least one additional pepsin cleavage sites are site is introduced by substituting one or more aspartic acid residues residue with one or more glutamic acid residues residue, substituting one or more tryptophan residues residue with one or more phenylalanine residues residue, and substituting one or more valine or one or more isoleucine residues residue with one or more leucine residues.
- 21. (Currently amended) The method as claimed in claim 20, characterized in that of Claim 20, wherein the degree of substitution which said modified Cry protein possesses is less than or equal to 25% of all aspartic acid, tryptophan, valine or isolecine residues.
- 22. (Currently modified) A polynucleotide encoding a the modified Cry protein as claimed in one of claims 1 to 11 of Claim 1.
- 23. (Currently amended) A chimeric gene comprising, functionally operatively linked to one another, at least:
 - a. one a promoter which is functional in a host organism;
 - b. a the polynucleotide as claimed in claim of Claim 22; and
 - c. a terminator element which is functional in a host organism.
- 24. (Currently amended) The chimeric gene as claimed in claim 23, characterized in that of Claim 23, wherein the promoter and the terminator element are functional in plants.
 - 25. (Currently amended) An expression or transformation vector containing a-the

chimeric gene as claimed in either of claims 23 and 24 of Claim 23.

- 26. (Currently amended) The vector as claimed in claim 27, characterized in that it is of Claim 25, wherein the vector is selected from the group consisting of a plasmid, a phage or and a virus.
- 27. (Currently amended) A host organism transformed with one of the vectors as claimed in either of claims 25 and 26 the vector of Claim 25.
- 28. (Currently amended) The <u>transformed</u> host organism as claimed in claim 27, eharacterized in that it is of Claim 27, wherein said transformed host organism is a plant.
- 29. (Currently amended) The plant as claimed in claim 28, characterized in that it contains, in addition to a chimeric gene as claimed in either of claims 23 and 24, of Claim 28, further comprising at least one other chimeric gene containing comprising a polynucleotide encoding a protein of interest.
- 30. (Currently amended) A part of a plant as claimed in claim 29 the plant of Claim 28.
- 31. (Currently amended) A seed Seeds from a the plant as claimed in claim 29 of Claim 28, wherein the seeds comprise the chimeric gene.
- 32. (Currently amended) A method for producing the modified Cry proteins as elaimed in one of claims 1 to 11, characterized in that it comprises at least the steps of a Cry protein modified to increase its sensitivity to degradation by pepsin comprising:
- a. culturing a the transformed host organism according to the invention of Claim 27 in a culture medium suitable for the growth and for the multiplication of said organism; and

- b. extracting the Cry proteins protein produced by the transformed host organism cultured in step (a).
- 33. (Currently amended) The method as claimed in claim 32, characterized in that it comprises a step (c) of purification of of Claim 32, further comprising purifying the Cry protein extracted in step (b).
- 34. (Currently amended) The method as claimed in either of claims 32 and 33, eharacterized in that the of Claim 32, wherein the transformed host organism is a microorganism.
- 35. (Currently amended) The method as claimed in claim 34, characterized in that the host of Claim 34, wherein the transformed host organism is a *Bacillus thuringiensis* bacterium.
- 36. (Currently amended) A monoclonal or polyclonal antibody, characterized in that it is directed against a modified Cry protein as claimed in one of claims 1 to 11 which specifically binds to the Cry protein of Claim 1.
 - 37. (New) The plant of Claim 29, wherein the plant is a monocotyledon.
- 38. (New) The plant of Claim 37, wherein the monocotyledon is selected from the group consisting of a corn plant, a wheat plant, and a rice plant.
 - 39. (New) The plant of Claim 29, wherein the plant is a dicotyledon.
- 40. (New) The plant of Claim 39, wherein the dicotyledon is selected from the group consisting of a rapeseed plant, a soybean plant, a tobacco plant and a cotton plant.